

Clinical trials on continuous measurement of fetal tcPco₂

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We have used the tcPco₂ registration in addition to the supervision of the fetus during labor by cardiotocography (CTG) and fetal blood analysis. We use a modified Severinghaus electrode (Radiometer E 5230) at a measuring temperature of 39°C respectively 44°C (2). By comparing the transcutaneously measured data with the values measured in the fetal blood it is possible to answer some specific questions of tcPco₂ monitoring in the fetus (1): In order to test the reliability of the tcPco₂ registration at different measuring temperatures we analysed the correlation between Pco₂ of the FBA and the synchronously registered tcPco₂ at 39°C ($r = 0.74$, $p < 0.001$, slope: 1.05, intercept: +8.57, $n = 111$) and compared this result with the analysis at 44°C (Fig. 1).

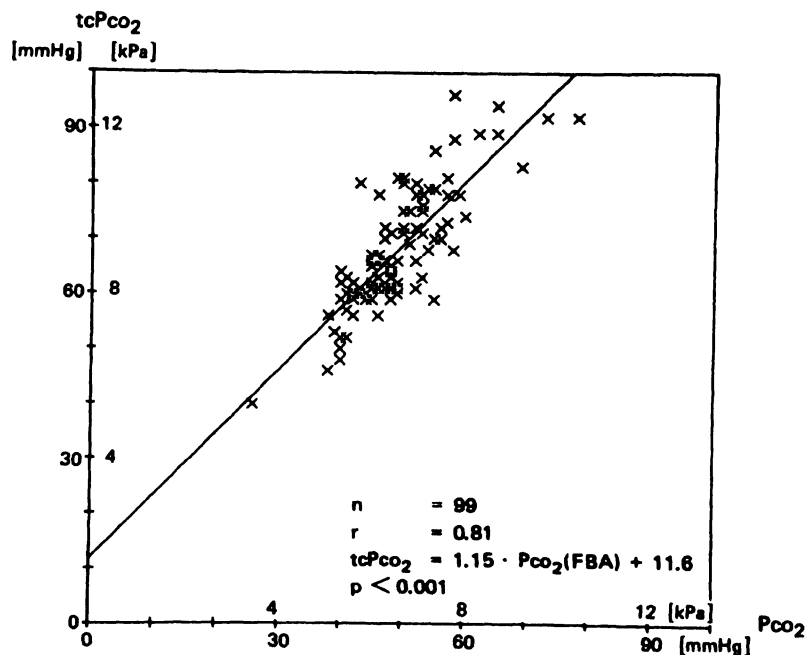


Fig. 1

While at both temperatures the correlation is significant ($p < 0.001$), due to Co₂ production in the tissue and the effect of the raised temperature, the transcutaneous values exceed the values of the fetal blood (at 44°C to a greater extent than at 39°C) (3). By comparing data of cases in which a caput succedaneum developed with cases where no such phenomenon occurred, we were able to elicit its influence on the tcPco₂ measurement (Tab. I).² We also analysed the influence of the duration of the transcutaneous measurement on its accuracy (Tab. II).

| caput succedaneum | correlation (R) | significance | intercept | slope | number of values |
|-------------------|-----------------|--------------|-----------|-------|------------------|
| 39°C - | 0.78 | < 0.001 | 10.84 | 1.01 | 28 |
| 39°C + | 0.72 | < 0.001 | 11.78 | 0.99 | 83 |
| 44°C - | 0.82 | < 0.001 | 14.68 | 1.09 | 74 |
| 44°C + | 0.80 | < 0.001 | 3.18 | 1.28 | 25 |

Tab. I

measurements at 44°C seem to be less accurate after registration of 150 min. As one basis of fetal surveillance is the intermittent pH measurement the correlation of the tcPco₂

A caput succedaneum has a considerable influence on the reliability of the tcPco₂ measurement especially when the prechosen temperature is 39°C while

| duration of measurement | correlation (R) | significance | intercept | slope | number of values |
|-------------------------|-----------------|--------------|-----------|-------|------------------|
| < 150min 39°C | 0.86 | < 0.001 | 11.92 | 0.98 | 13 |
| > 150min 39°C | 0.83 | < 0.001 | 7.60 | 1.11 | 14 |
| < 150min 44°C | 0.93 | < 0.001 | 8.16 | 1.21 | 24 |
| > 150min 44°C | 0.79 | < 0.001 | 7.41 | 1.24 | 20 |

Tab. II

with the pH is of great clinical importance. In the range between a pH level of 7.25 to 7.50 we find the tcPco₂ values to be rather scattered. However with pH value dropping to 7.24 (pre-acidosis), all

tcPco₂ values of this study at 44°C and not corrected, are >70 mmHg (Fig. 2).

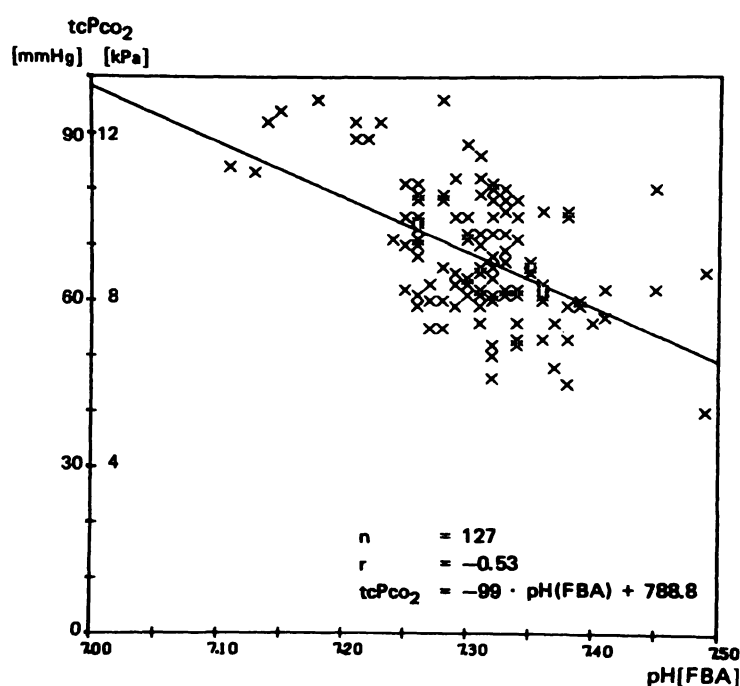


Fig. 2

If we only then perform an FBA when the indication emerged from an irregular heart rate pattern, and additionally this level of 70 mmHg is exceeded, a considerable number of FBAs could be spared. (This action line should be confirmed by additional measurements). Furthermore the timing of an obstetrical operation may be optimised. During our clinical trial the occurrence of an acute complication was in all instances clearly indicated by a rise of the tcPco₂ value from its original level. All these babies, in spite of acidosis, were born by immediate operation before clinical signs

of depression (Apgar<7) had manifested.

Bibliography

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